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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/695,994	10/30/2003	Lopa Mishra	P04470US02/BAS	7531
881 75	90 04/20/2006		EXAMINER .	
STITES & HARBISON PLLC			MERTZ, PREMA MARIA	
1199 NORTH FAIRFAX STREET SUITE 900			ART UNIT	PAPER NUMBER
ALEXANDRIA, VA 22314			1646	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)
	10/695,994	MISHRA, LOPA
Office Action Summary	Examiner	Art Unit
	Prema M. Mertz	1646
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with the c	orrespondence address
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING D. Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. If NO period for reply is specified above, the maximum statutory period v. Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim will apply and will expire SIX (6) MONTHS from , cause the application to become ABANDONE	I. lely filed the mailing date of this communication. (35 U.S.C. § 133).
Status		
1) Responsive to communication(s) filed on 2 2 2a) This action is FINAL. 2b) This 3) Since this application is in condition for alloware closed in accordance with the practice under E	s action is non-final. nce except for formal matters, pro	
Disposition of Claims		
4) ⊠ Claim(s) 1-18,20 and 21 is/are pending in the state 4a) Of the above claim(s) 1-18 and 20 is/are with 5) □ Claim(s) is/are allowed. 6) ⊠ Claim(s) 21 is/are rejected. 7) □ Claim(s) is/are objected to. 8) □ Claim(s) are subject to restriction and/or	ithdrawn from consideration.	
Application Papers		
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) acc Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Examine 11.	epted or b) objected to by the Bedrawing(s) be held in abeyance. See ition is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		•
12) ☐ Acknowledgment is made of a claim for foreign a) ☐ All b) ☐ Some * c) ☐ None of: 1. ☐ Certified copies of the priority document 2. ☐ Certified copies of the priority document 3. ☐ Copies of the certified copies of the priority document application from the International Bureau * See the attached detailed Office action for a list	s have been received. s have been received in Application rity documents have been received u (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	4) N Interview Summary Paper No(s)/Mail Da	ite
3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date	6) Other:	atent Application (PTO-152)

Claims 1-18, 20-21 are pending in the instant application.

Election/Restriction

1. Applicants election with traverse of antibodies to an elf-3 protein of amino acid sequence set forth in SEQ ID NO:7 is acknowledged. The traversal is on the ground(s) that the restriction is improper since the examiner has not shown that examination of antibodies to elf, praja-1 and pk proteins, would entail a serious burden. This argument is not found persuasive because the Examiner has cited art to demonstrate that the Groups drawn to different antibodies to the disparate proteins lack a special technical feature that is unique and is absent from the prior art.

Each of the antibodies to elf, praja-1 and pk proteins does not share a common technical feature, which is based on a common property or special technical feature not found in the prior art. These antibodies are independent and distinct chemical compounds lacking either a common structural property which distinguishes them as a group from structurally related compounds of the prior art or which provides them with a common utility, which is lacking from those prior art elements.

The first claimed invention fails to recite such a feature, since claim 21 recites an antibody to the amino acid sequence of SEQ IDNO:7, which antibody is found in the prior art.

The claim encompasses any antibody which can bind to any epitope in the amino acid sequence as set forth in SEQ ID NO:7 and, therefore the claim encompasses an antibody to dystrophin (see WO 89/06286), which has 11 amino acids in common with the elf-3 protein of the instant invention. The reference discloses the antibody (page 6, lines 7-10; pages 27-29) meeting the limitations of an antibody that binds to a polypeptide of the amino acid sequence of SEQ ID

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NO:7. Since the first claimed invention lacks a special technical feature, the other claimed inventions cannot share a special technical feature with the first claimed invention.

Having shown that these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their recognized divergent subject matter as defined by MPEP 808.02, the Examiner has prima facie shown a serious burden of search (see MPEP 803). Therefore, an initial requirement of restriction for examination purposes as indicated is proper. Therefore, Groups drawn to antibodies to elf, praja-1 and pk proteins lack a special technical feature that is unique and absent from the prior art and the restriction requirement set forth on 11/21/2005 is proper and is being maintained.

The Groups as delineated in the restriction requirement (11/21/2005) meet the requirements to support a restriction between the Groups.

The requirement is still deemed proper and is therefore made FINAL.

Claim 1-18, 20 are withdrawn from further consideration by the examiner, as being drawn to a non-elected invention.

Specification

- 2a. The title of the invention is not descriptive. A new title is required that is clearly indicative of the invention to which the claims are directed. It is suggested that the title be amended to recite "antibodies to elf-3 protein".
- 2b. Applicants are requested to update the status of the prior applications to which the instant application is claiming benefit. The status of nonprovisional parent 08/841,349 should be updated and the expression, "Patent No. 5,955,594" should follow the filing date of the parent

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application. Furthermore, the status of nonprovisional parent 09/431,184 should be updated and the expression, "Patent No. 6,642,362" should follow the filing date of the parent application.

Claim rejections-35 USC § 112, second paragraph

3. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claim 21 is rejected under 35 U.S.C. § 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 21 is vague and indefinite because it recites "binds to an amino acid sequence as set forth in SEQ ID NO:7" rather than "binds to a protein consisting of the amino acid sequence as set forth in SEQ ID NO:7", because the specific amino acid sequence is a characteristic or property of the protein.

Claim Rejections - 35 USC § 102

4. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claim 21 is rejected under 35 U.S.C. 102(b) as being anticipated by WO 89/06286.

The reference discloses an antibody to dystrophin (see page 6, lines 7-10; pages 27-29) which has 11 amino acids 100% identical with elf-3 protein amino acid sequence set forth in SEQ ID NO:7 (see Sequence Comparison 'A' attached). The antibody of the reference meets the

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limitations of an antibody that binds to an amino acid sequence as set forth in SEQ ID NO:7.

Therefore, the antibody of the reference anticipates instant claim 21.

Conclusion

No claim is allowable.

Claim 21 is rejected.

Advisory Information

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Prema Mertz whose telephone number is (571) 272-0876. The examiner can normally be reached on Monday-Friday from 7:00AM to 3:30PM (Eastern time).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Janet Andres, can be reached on (571) 272-0867.

Official papers filed by fax should be directed to (571) 273-8300. Faxed draft or informal communications with the examiner should be directed to (571) 273-0876.

Information regarding the status of an application may be obtained from the Patent application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Prema Mertz Ph.D., J.D. Primary Examiner Art Unit 1646 March 29, 2006 ö

Gaps

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Mismatches

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11; Conservative

Matches

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providing improved plant growth and development under at least one stress condition, improved lignin production or improved galactomannan production. This sequence represents a bacterial polypeptide used in the scope of the invention. Note: The sequence data for this patent did not form part of the printed specification but was obtained in electronic format from USPTO at seqdata.uspto.gov/sequence.html.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        New nucleic acid encoding PRO polypeptide, useful for diagnosing and treating an immune related disorder, e.g. systemic lupus erythematosus, rheumatoid arthritis, osteoarthritis, autoimmune hemolytic anemia, or
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Immunosuppressive; Antirheumatic; Antiarthritic; Osteopathic;
Muscular-Gen.; Vasotropic; Antianemic; Antipsoriatic; Immunostimulant.
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0; Mismatches 0; Indels
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Best Local Similarity 100.v.
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diagnosis; prenatal;
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heterozygote; gene therapy; genetic screening; foetal screening
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                                                                                                                                                       Sequence encoded by human muscular dystrophy (MD) cDNA.
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deposited as ATCC 58666-57677. The MD
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42 VOKKTFTKWVN 52
                    70 VOKKTFTKWW 80
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10-JAN-1990
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myopathies, Sjögren's syndrome, systemic vasculitis, sarcoidesis, autoimmune hemolytic anemia, autoimmune or immune-mediated skin diseases including bullous skin diseases, erythema multiforme and contact dermatitis, psoriasis, lymphadenopathy, splenomegaly and leukopenia. The present sequence represents the amino acid sequence of a human PRO polypeptide.

Length 2000;

DB 9; 0.29;

5.0%; Score 11; 100.0%; Pred. No.

Best Local Similarity

Query Match

Sequence 2000 AA;

compound or composition, and methods are useful for diagnosing and treating an immune related disorder, e.g. systemic lupus erythematosus, rheumatoid arthritis, osteoarthritis, juvenile chronic arthritis,

invention relates to an isolated nucleic acid. The polypeptide

spondyloarthropathies, systemic sclerosis, idiopathic inflammatory